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The sicca syndrome in thalassaemia major

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Abstract

A 20 year old man with β thalassaemia developed symptoms of the sicca syndrome. His serum contained rheumatoid factor and antinuclear antibodies. A biopsy specimen of labial salivary gland showed large accumulations of haemosiderin within the parenchymal cells of the acini.

Although in this case the sicca syndrome could not be definitely distinguished from Sjögren's syndrome, the patient's HLA type was not the one usually associated with Sjögren's syndrome. Histological appearances suggested that the causative factor of the sicca syndrome was iron overload owing to an intensive blood transfusion regimen.

Introduction

Iron overload has been reported to have produced the sicca syndrome, xerostomia, and xerophthalmia in a patient with idiopathic haemochromatosis.¹ We report here the first case of the sicca syndrome in a young adult with β thalassaemia major and severe iron overload owing to repeated blood transfusions.

Case report

The patient was a 20 year old man in whom β thalassaemia major had been diagnosed at the age of 1 year. Transfusions were started a month later and continued up to the time of writing at intervals of one to two months. The mean haemoglobin concentration before transfusion had not been below 9.3 g/dl throughout the past three years. We calculated from clinical records that he had received a total of over 100 l blood, containing about 50 g iron. Splenectomy was performed at age 3. There was no history of viral hepatitis. Chelation treatment was started in 1975 with desferrioxamine mesylate intramuscularly, 500 mg/day five days a week; from the beginning of 1980 he self administered an average dose of 2 g desferrioxamine mesylate subcutaneously over eight hours on alternate days. In the past year, in an attempt to reduce the iron load, he had undergone

monthly wash outs with intravenous desferrioxamine mesylate, 300 mg/kg/24 h (16 g/24 h). In 1981, at age 19, he developed diabetes requiring insulin treatment. In July 1982 he had an episode of cardiac failure, which responded readily to diuretics and digitalis. Puberty did not develop spontaneously, and he was therefore given human chorionic gonadotrophin with some improvement in sexual maturation.

In April 1983 he complained of dry mouth and itchy eyes of a few weeks' duration. Pertinent clinical findings, apart from a dry scaly tongue, included moderately hyperpigmented skin and a large, hard liver palpable 7 cm below the right costal margin. No swelling of the parotid or lachrymal gland was noticeable. Liver function tests showed an alanine transferase activity three times the upper limit of normal, prothrombin activity 50% of normal, albumin concentration 35 g/l, and gammaglobulin concentration 3.2 g/100 ml. Serum was negative for hepatitis B surface antigen and anti-e antibodies and positive for anti-s and anti-c antibodies. Fibrinogen concentration was 2.8 g/l. Rheumatoid factor was present in the serum, and antinuclear antibodies were present at a titre of 1/500, with a speckled pattern. No organ specific antibodies were found. HLA typing showed B13, Bw35, DR5, and DR7. The serum ferritin concentration was 3.3 μ g/l, and the average urinary excretion of iron after subcutaneous desferrioxamine mesylate was 645 μ mol (36 mg)/24 h; excretion doubled when the high intravenous dose was administered. Faecal excretion of chymotrypsin was below normal. Schirmer's test gave a result of 3 mm.

Ophthalmological examination with fluorescein showed areas of abnormal epithelial cells on the cornea. A biopsy specimen of the labial salivary gland showed large accumulations of haemosiderin within the parenchymal cells of the acini. There was no evidence of lymphocytic infiltrate, cellular destruction, or interstitial fibrosis (figure).

Treatment with bromhexine and artificial tears was prescribed with little subjective improvement.

Specimen of labial gland. The salivary acini are filled with haemosiderin granules. (Perl's stain. $\times 43$.)

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Comment

Most pathological manifestations of thalassaemia major are attributed to haemosiderosis owing to transfusions. Iron deposits in the heart, liver, pancreas, and endocrine glands have been described.² In this patient with the sicca syndrome we found heavy deposition of iron in the labial salivary glands similar to that described in a woman with idiopathic haemochromatosis.¹ We were unable to distinguish the syndrome from Sjögren's syndrome on clinical grounds. Sjögren's syndrome is an autoimmune disorder affecting mainly postmenopausal women. It is the result of lymphocyte mediated destruction of the exocrine glands and can occur alone or in association with another connective tissue disorder, most commonly rheumatoid arthritis. Our patient never had symptoms of connective tissue disease or enlargement of the parotid gland. His HLA type was not the one (B8 and DR3 or 4) usually associated with Sjögren's syndrome. Thyroid autoantibodies were not present. He did have hypergammaglobulinaemia and was positive for rheumatoid factor and antinuclear antibodies. High concentrations of gammaglobulin, however, are almost invariably present in patients with thalassaemia,³ and rheumatoid factor and antinuclear antibodies have been found in 31% and 13% respectively of a series of unselected patients with β thalassaemia major⁴ and have been attributed to the repeated antigenic stimuli received with transfusions. The histological appearance of the salivary glands, with heavy deposition of iron in glandular acini and absence of lymphocytic infiltration, did not support a diagnosis of Sjögren's syndrome but suggested damage related to iron.

Although we could not with certainty rule out associated autoimmune disease, we are inclined to regard the sicca syndrome in

our patient as a consequence of iron overload. Disease of the exocrine portion of the pancreas leading to reduced serum chymotrypsin concentrations has been described in patients with thalassaemia and has been attributed to infiltration of the acinar tissue by iron.⁵ With the improvement in life expectancy due to intensive transfusion regimens and aggressive chelation of iron, doctors taking care of patients with thalassaemias are likely to see new symptoms developing in this previously non-existent population of adult patients. The sicca syndrome may be one of them.

Professor G Veneroni, Centro Trasfusionale, Ospedale Fatebenefratelli, performed the HLA typing.

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Glycosylation of hair: possible measure of chronic hyperglycaemia

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Abstract

To determine whether hair is excessively glycosylated in diabetes mellitus 4 cm hair samples were taken proximally from behind the ear in 50 white non-diabetics and 46 diabetics. Hair glycosylation was assayed by a modification of the thiobarbituric acid reaction. Blood was taken from the diabetics at the same time for measurement of glycosylated haemoglobin concentration.

The mean (1 SD) concentration of fructosamine ($\mu\text{mol}/100 \text{ mg hair}$) was 0.054 (0.011) for normal hair. Glycosylation was not related to sex, age, or hair colour. The diabetics' hair was more heavily glycosylated (0.097 (0.045)) than normal ($p < 0.01$) and there was a

correlation between hair glycosylation and the concentration of glycosylated haemoglobin in the diabetics ($r = 0.71$; $p < 0.01$). Hair from non-diabetics showed a stable time related increase in glycosylation when incubated with glucose.

Glycosylation of hair might provide a stable long term measure of tissue glycosylation, useful in the investigation of microvascular complications of diabetes mellitus.

Introduction

The discovery of increased non-enzymatically glycosylated haemoglobin concentrations in diabetes mellitus¹⁻⁴ has led to intensive research into similar excess glycosylation of other tissue proteins, especially in an attempt to establish a link between this process and chronic complications of the condition. Measurement of the percentage concentration of glycosylated haemoglobin has proved useful in assessing diabetic control, but tissue collagen, which is susceptible to functional changes from excessive glycosylation, has a much slower rate of turnover than haemoglobin.⁵ It is therefore of interest to investigate possible longer term markers of hyperglycaemia than haemoglobin, and we have conducted such a study.

As it is not feasible repeatedly to sample tissues containing basement membrane from the same subject, we investigated the

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